

# SYNTHESIS AND PHOTOPHYSICAL STUDY OF ALQ<sub>3</sub>: P ORGANIC PHOSPHORS

Sunil A. Bhagat\*

Department of Physics, Kamla Nehru Mahavidyalaya, Nagpur, 440024, India

## ABSTRACT

**Phosphorus** doped alumium-8hydroxyquinoline metal complex is synthesized using simple precipitation route in order to reduce the cost and time of material. It is characterized by X-ray Diffractogram of P doped Alq<sub>3</sub> complex exhibits well defined X-ray diffraction lines, confirming its crystalline nature. Photo luminescence spectra of Alq<sub>3</sub>: P reveals intensity in green region. The synthesized P doped Alq<sub>3</sub> organic phosphor is promising material for optical or optoelectronic applications.

Keywords:organic phosphor, Alq<sub>3</sub>, Xrd, Photoluminescence, P doped Alq<sub>3</sub>

## 1. Introduction

Now davs Organic light-emitting diodes (OLEDs) constitute a rapidly developing field. Many believe that they represent the future of flat panel display technology. The main focus of OLED research is to address these issues. Two types of compounds: conjugated organic poly polymers, such as (1,4phenylenevinylene) (PPV) and molecular species such as aluminum tris-(8hydroxyquinolate) (Alq<sub>3</sub>) are used frequently. Organic have attracted tremendous interests for application in functional nanoscale electronic and optoelectronic devices, the crystalline and molecular arrangement of which have a great influence on the performance of these devices. compared [1-4] However. with the overwhelming majority of inorganic nonmaterial, only a few successful preparations of organic one-dimensional (1D) nonmaterial such as nanowires and nanotubes are reported. [3, 5-7]

Since the first efficient low-voltagedriven organic light-emitting diodes (OLEDs) based on tris (8-hydroxyquinoline) aluminum (Alq<sub>3</sub>) were reported [8]. Alq<sub>3</sub> has become a important prototypical electron transport and emitting material for OLED devices because of its excellent stability and electro-luminescence properties. Therefore lumophores based on metallo-8-hydroxyquinolate Aluminum prepared from wet Chemical method and codoped with Phosphorus non metal element are prepared and characterization of the material by X-Ray diffraction spectrographic is carried out. Alumium-8-hydroxyquinoline and co-doped materials with varying concentrations of dopants [9, 10, 11, 12] are synthesized by simple wet Chemical route in order to reduce the cost and time of synthesized material. In the synthesis technique when Phosphorus non metal is added which contribute an electron. withdrawing constituent at the 5-position in 8hydroxyquinoline, increasing the solubility of the corresponding metal quinolate complexes in non polar solvents.

# 2. Experimental

Alq<sub>3</sub> was prepared as follows: firstly take 25 ml double distilled water and 25 ml acetic acid in beaker. Dissolve 5 gm of 8hydroxyquinoline in a mixture of double distilled water, acetic acid and stir it still the orange transparent solution was obtained say solution I.Take 5gm Al (NO<sub>3</sub>)<sub>3</sub>.9H<sub>2</sub>O and dissolve in double distilled water. Stir it till clear solution was obtained say solution II. Mix the solution I and II and stir for 10 min and add N<sub>4</sub>OH solution by drop by drop to this mixture of solution with continuous stirring. Filter the vellow green precipitate and wash the precipitate with double distilled water for 8 to 10 times. Place the precipitate for drying 40-50°C. The other derivative of 8hydroxyquinoline metal complex is prepared by Simple precipitation method same as Alq<sub>3</sub>. Aluminium nitrate is replaced by ammonium dihydrogen phosphate (NH4)H<sub>2</sub>PO<sub>4</sub>. For the preparation of  $Alq_3$  host lattice, aluminium nitrate and 8-hydroxyquinoline as raw materials and for  $Alq_3$ : P, ammonium dihydrogen phosphate is used as other material to dope phosphorus with proportion 5%,3%,2% and 1% mixed in an appropriate molar ratio mentioned in equation 1.

# 3. Results & Discussion

# 3.1. XRD OF Alq<sub>3</sub>: P

Diffraction data has historically provided regarding information the structures of crystalline solids. Such data can be used to determine molecular structures, ranging from Simple to complex, since the relative atomic positions of atoms can be determined. X-ray Diffraction provides important evidence and indirect proof of atoms. The Symmetry of the diffraction patterns corresponds to the symmetry of the atomic packing. It is the simplest way to determine inter atomic lattice spacing that exists. The intensity of the depends diffracted beams also on the arrangement and atomic number of the atoms in the repeating motif, called the unit cell. Thus, the intensities of diffracted spots calculated for trial atomic positions can be compared with the experimental diffraction intensities to obtain the positions of the atoms themselves. The XRD pattern did not indicate presence of the constituents like nitrates, ammonia and other likely phases. This result indicates the final product formed is in crystalline and homogeneous form.

The synthesized complex have been characterized by XRD on the 'Expert pro' Automated power Diffractometer system company name Analytical, Netherland taken at 'SAIF' Punjab University, Chandigarh as shown in Fig.1.

## 3.2. PL properties of Alq<sub>3</sub> and Alq<sub>3</sub>: P

Excitation and emission spectra exhibit Photo luminescence properties of Alq<sub>3</sub> powder by excitation wavelength is 384 and 455 nm. The prominent PL emission peak is observed at 531 nm in green region of the spectrum well matched with the green emission of yellowish green powder earlier reported attributed to  $\alpha$ phase of meridianal isomer of Alq<sub>3</sub>. The PL

excitation spectrum at 455nm of prepared Alg<sub>3</sub> powder reveals that prepared phosphors is not only suitable for organic light emitting diode but also for photoluminescence liquid crystal display (PLLCD) and solid state lighting applications. See Fig. 2(a) and (b). Jablonski diagram illustrating possible electronic processes following absorption of a photon for Alq<sub>3</sub>  $K_r$  and  $K_{nr}$  indicate the radiative and nonradiative transitions and here only singlet to singlet  $(S_1, S_0)$  is radiative and remaining transitions are non-radiative. See Fig. 3(a). This electronic transition is supported by adiabatic potential model of dynamical relaxation process for electroluminescence in Alq<sub>3</sub> reported by Akai et.al.  $S_1 S_0$  See Fig. 3(b).

Photo luminescence (PL) excitation spectrum of Alq<sub>3</sub>: P having excitation wavelength is 440 nm exhibits the prominent PL excitation peak of Alq<sub>3</sub>: P is observed at 438 nm See Fig. 4(a). PL emission spectrum of prepared Alq<sub>3</sub>: P powder shows the prominent shoulder at 500nm Alq<sub>3</sub>: P (3%) with higher intensity, 500nm Alq<sub>3</sub>: P (5 %), 502nm Alq<sub>3</sub>: P (1%), and 502nm Alq<sub>3</sub>: P (2%) with decreasing intensity in green region. See Fig. 4(b) Hence the emission peak reveals that prepared phosphors is not only suitable for organic light emitting diode but also for photoluminescence liquid crystal display (PLLCD) and solid state lighting application as their no significant change in the emission peaks but there is an increase in the intensity. See table 1.

## 4. Conclusion

Alq<sub>3</sub>: P hybrid organic phosphor is synthesized by the precipitation wet chemical method. Phosphor is characterized by XRD spectrograph. X-ray spectrograph of P doped Alq<sub>3</sub> complex displays well defined X-ray diffraction lines, confirming its crystalline nature and its grain size. Photo luminescence excitation spectra observed at 440nm violet region whereas emission spectra reveals at 500nm in the green region for combination Alq<sub>3</sub>: P (3%). So Alq<sub>3</sub>: P organic phosphor is suitable in the green region for PLLCD and OLED, nano rods, nano wires and solid state lighting application devices.

1.1	. Table	e 1					
					Emission		
		Organic	Excitation	Emission	Intensity		
		phosphors	Wavelength(nm)	Wavelength(nm	) (arb.		
			201 nm	521 mm	<u>unit)</u>		
		Alq <sub>3</sub>	384 IIII 455 nm	531 mm	241 450		
		$A10 \cdot P$	433 IIII 438 nm	500nm	430		
		(0.5%)	450 1111	5001111	233		
		$AlO_3$ : P	438 nm	500 nm	305		
		(3%)		2000 1111	000		
		AlQ <sub>3</sub> : P	433nm	502 nm	112		
		(1%)					
		AlQ <sub>3</sub> : P	436 nm	502nm	85		
		(2%)					
		Counts					
		Alq3 P 0.5	%				
		10000 —					
		5000 -					
		0000					
			M M Marra				
		0	20 30 40	50 60 70	80 90	<u></u>	
			$\mathbf{E}_{\mathbf{z}} = 1 \cdot \mathbf{V} \mathbf{I}$	Position [°2Theta] (Copper (Cu))			
	500		F1g. 1: Af	$500 \neg$			
ensity(arb. unit)	500 ]				b		
		•	$\cdot \cdot \Lambda$	400	$\langle \neg \rangle$		
	400 -		/ \	£ 400 1	- /		
				5			
	300 -			<u>e</u> 300 -	( )		
			$\gamma$	y(a		۱.	
	200 -	and the second second	Ŭ l	- 200 -	$1/$ $\times$	\	
nte		prend		ter	$ I  = \langle$	1	
_	100 -	/		드 100 -	// `	$\langle \rangle$	
		,	ι	100	1/		
	0				/		
	200	300	400 500	400	500	600	700
	Wavelength(nm)			400	ουυ ∖N/aγelenαth	(nm)	700
	(i)				(ii)		

Fig 2: (i) Excitation and (ii) Emission spectra of  $Alq_3$ 





Fig. 4- (a) Excitation spectrum: Alq3: P (b) Emission spectrum: Alq<sub>3</sub>: P Equation

 $\begin{array}{rcl} A1 (NO3)3.9H2O + (C9H7NO) & \rightarrow & A1 (C9H7NO) 3 + 9H2O + 3(NO2) \uparrow \\ A1 (NO3)39H2O + (NH4) H2PO4 + 4(C9H7NO) & \rightarrow & A1 P (C9H7)O4 + 9H2O + NH3 + HNO3 \end{array}$ 

#### REFERENCES

[1] Veinot J. G. C., T. J. Marks, Acc. Chem. Res., 2005;38; 632.

[2] Chiu J. J., Kei C. C., Perng T. P. & Wang W. S., Adv. Mater., 2003;15; 1361.

[3] Liu H. B., Zhao Q., Li Y. L., Liu Y., Lu F.
S., Zhuang J. P., Wang,S. Jiang L., Zhu D. B.,
Yu D. P. & Chi L. F., J. Am. Chem. Soc.,
2005;127; 1120.

[4] Kastler M., Pisula W., Laquai F., Kumar A., Davies R.J., Baluschev S., Garcia-Gutie 'rrez M. C., Wasserfallen D. ,H. Butt J., Riekel C., Wegner G., & Mullen K. , Adv. Mater., 2006; 18; 2255.

[5] Fu H. B., Xiao D. B., Yao J. N., Yang G.Q. & Angew, Chem., Int.Ed., 2003;42; 2883.

[6] Hu J. S., Guo Y. G., Liang H. P., Wan L. J.& Jiang L., J. Am. Chem.Soc., 2005; 127; 17090.

[7] Lee J. K., W. Koh K., W. Chae S. & Y. Kim R., Chem. Commun.,2002; 138.

[8] Tang C. W. & Van Slyke S. A., J. Appl. Phys., 1989; 3610.

[9] Bhagat S.A., Raut S.B., & Dhoble S.J., J. Bio. Chem. Lumin, 2012; 27(4); 2431

[10] Bhagat S.A., Borghate S.V., Thejo Kalyani N, Dhoble S.J., J. Bio. Chem. Lumin, 2013; 2; 433-439.

[11] Bhagat,S.A. Borghate S. V., Thejo Kalyani N, Dhoble S.J., J. Bio. Chem. Lumin, 2015; 30(3), 251-256.

[12] Bhagat S.A., Borghate S.V., Koche N.S., Thejo Kalyani N, Dhoble S.J., J. OPTIK, 2014;125(15);433-439.